

Page 5, please insert immediately before the line which read "Anti-mitotic Activity *In Situ*" the following new header:

DETAILED DESCRIPTION OF THE INVENTION

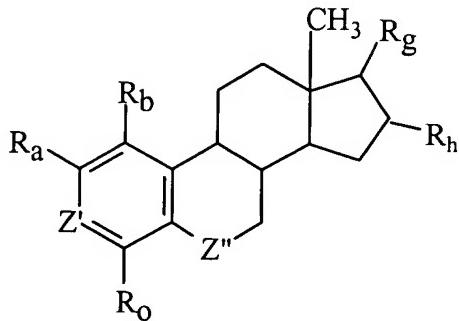
Please rewrite page 10, last paragraph, as follows:

It should be understood that in addition to the ingredients, particularly mentioned above, the formulations of this invention may include other agents conventional in the art having regard to the type of formulation in question, for example, those suitable for oral administration may include flavoring agents.

In the Claims

Please cancel Claims 1-40 without prejudice and enter the following new claims.

Rule 126 26 -41: (New) A compound of the formula:



wherein:

- a) R_a is $-OR_1$ or $-OCOR_1$, wherein R_1 is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle;

b) R_b and R_o are independently selected from -H, -Cl, -Br, -I, -F, -CN, -OH, aryl, aralkyl, alkenyl, alkynyl, heterocycle, $-(CH_2)_nOH$ where n is from 1 to 6, straight or branched alkyl with up to 10 carbons, substituted alkyl with up to 10 carbons; $N(R_2)(R_3)$, -OR₂, or -OCOR₂, wherein R₂ and R₃ are independently selected from H, alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aryl, aralkyl, alkenyl, alkynyl, or heterocycle;

c) Z' is >CH; >COH; >CR₄OH, where R₄ is an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, or where R₄ is aralkyl, aryl, alkenyl, alkynyl, or heterocycle;

d) >C-R_g and C-R_h are independently selected from >CH₂, >CHR₅, >CR₅R₆, >C(H)-OH, >C=O, >C=N-OH, >C(R₅)OH, >C=N-OR₅, >C(H)-NH₂, >C(H)-NHR₅, >C(H)-NR₅R₆, or >C(H)-C(O)-R₅, or >C(R₅)-C(O)R₆ where each R₅ and R₆ is independently selected from an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aralkyl, alkenyl, alkynyl, or heterocycle; and

e) Z" is >CH₂, >C=O, >C(H)-OH, >C=N-OH, >C=N-OR₇, C(H)-C≡N, or >C(H)-NR₇R₈, wherein R₇ and R₈ are independently selected from H, an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aralkyl, alkenyl, alkynyl, or heterocycle;

and wherein the compound is not 2-methoxyestradiol.

27

26

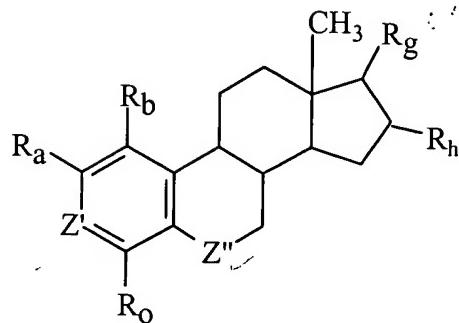
42. (New) The compound of Claim 41, wherein R_a is -OR₁, wherein R₁ is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle.

29

43. (New) The compound of Claim 41, wherein R_a is -OCOR₁, wherein R₁ is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle.

26

29
44. (New) A method of inhibiting neovascularization in a mammal, comprising administering to the mammal a neovascularization-inhibiting amount of a compound of the formula:



wherein:

- a) R_a is -OR₁ or -OCOR₁, wherein R₁ is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle;
- b) R_b and R_o are independently selected from -H, -Cl, -Br, -I, -F, -CN, -OH, aryl, aralkyl, alkenyl, alkynyl, heterocycle, -(CH₂)_nOH where n is from 1 to 6, straight or branched alkyl with up to 10 carbons, substituted alkyl with up to 10 carbons; N(R₂)(R₃), -OR₂,

or $-\text{OCOR}_2$, wherein R_2 and R_3 are independently selected from H, alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aryl, aralkyl, alkenyl, alkynyl, or heterocycle;

c) $\text{-Z}'$ is $>\text{CH}$; $>\text{COH}$; $>\text{CR}_4\text{OH}$, where R_4 is an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, or where R_4 is aralkyl, aryl, alkenyl, alkynyl, or heterocycle;

d) $>\text{C-R}_g$ and C-R_h are independently selected from $>\text{CH}_2$, $>\text{CHR}_5$, $>\text{CR}_5\text{R}_6$, $>\text{C(H)-OH}$, $>\text{C=O}$, $>\text{C=N-OH}$, $>\text{C(R}_5\text{)OH}$, $>\text{C=N-OR}_5$, $>\text{C(H)-NH}_2$, $>\text{C(H)-NHR}_5$, $>\text{C(H)-NR}_5\text{R}_6$, or $>\text{C(H)-C(O)-R}_5$, or $>\text{C(R}_5\text{)-C(O)R}_6$ where each R_5 and R_6 is independently selected from an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aralkyl, alkenyl, alkynyl, or heterocycle; and

e) Z'' is $>\text{CH}_2$, $>\text{C=O}$, $>\text{C(H)-OH}$, $>\text{C=N-OH}$, $>\text{C=N-OR}_7$, $\text{C(H)-C}\equiv\text{N}$, or $>\text{C(H)-NR}_7\text{R}_8$, wherein R_7 and R_8 are independently selected from H, an alkyl, branched alkyl, or substituted alkyl with up to 10 carbons, aralkyl, alkenyl, alkynyl, or heterocycle.

SEARCHED INDEXED
SERIALIZED FILED
APR 22 1992
RECEIVED
U.S. PATENT AND TRADEMARK OFFICE

30

28

45. (New) The method of Claim 44, wherein R_a is $-\text{OR}_1$, wherein R_1 is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle.

30

28

46. (New) The method of Claim 44, wherein R_a is $-\text{OCOR}_1$, wherein R_1 is a straight, branched, or substituted alkyl with up to 10 carbons, aralkyl, aryl, alkenyl, alkynyl, or heterocycle.